Amendment to the Specification:

Please amend the specification at page 1 by replacing the last line with the following:

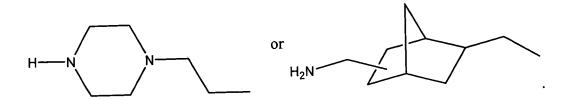
X is -O-, [[-COO-]] $\underline{-C(=O)O}$ or $-CHR_4$ -, with R_4 and R_3 together forming an ethylene group,

Please amend the specification at page 3 by replacing the formula for the tetravalent radical with the following formula:

with the following formation
$$H_3C$$
 CH_3 CH_2 CH_2

Please replace the sixth paragraph on page 4 of the specification with the following paragraph:

Special preference is given to compounds of formulae Ia and Ib wherein [[R₁]] \underline{R}_5 is n-butyl, n-octyl, cyclohexyl, benzyl, 2-aminoethyl, 4-(aminomethyl)pentyl, 5-amino-2-methylpentyl, 3-dimethylaminopropyl, 3-methylaminopropyl, 4-aminocyclohexyl or a radial of formula -CH₂CH₂NHCH₂CH₂NH₂,



Please amend the specification at page 5 by replacing the last paragraph with the following:

The episulfide of formula IIIa is then dissolved in an aprotic or protic organic solvent and, under inert gas (argon or nitrogen), reacted with the amine R_5 -NH- R_2 , the amount of the amine R_5 -NH- R_2 preferably being so selected that there are from 1 to 10 [[NH]] amine groups for one episulfide group. Preferred solvents are alcohols (e.g. methanol, ethanol, tertbutanol) and aromatic hydrocarbons, for example toluene and xylene

Please amend the specification at page 7 by replacing the first full paragraph with the following:

Episulfides can, for example, also be synthesized synthesized from the corresponding epoxides by reaction with triphenylphosphine sulfide.

Please replace the first full paragraph on page 11 with the following paragraph:

As component [[A]] C of the substance mixtures in accordance with the invention, preference is given to cycloaliphatic and aliphatic amines, especially the amines of formulae R₅-NH-R₂ and E-(NHR₂)_{m+1} used for preparation of the polymercaptopolyamines according to the invention.

Please replace the paragraph bridging pages 12 and 13 with the following paragraph:

- I. <u>Preparation of compounds of formula [[I]] Ia and Ib</u>
- a) General procedure for the preparation of polyepisulfides:

 The polyepoxide of formula II is epoxy compounds of formula IIa or IIb

 are dissolved in an amount of solvent that is from 0.5 to 5 times the

 amount of the polyepoxide epoxy compound and is stirred, under nitrogen,

with thiourea or alkali-metal or ammonium thiocyanate (0.8-1.2 equivalents of sulfur per epoxy equivalent) at 60-100°C until the epoxy content has fallen to nearly zero.

After separating off the by-products by means of filtration, extraction or phase separation, the polyepisulfide is isolated as a result of concentration by evaporating off the solvent.

b) General procedure for the preparation of polymercaptopolyamines: The polyepisulfide is dissolved in an amount of solvent that is from 0.5 to 5 times the amount of the polyepisulfide and under nitrogen and with vigorous stirring, is combined with the amine (either R₅-NH-R₂ and E-(NHR₂)_{m+1}), which likewise has been dissolved in an amount of solvent that is from 0.5 to 5 times the amount of the amine. The amount of amine is selected so that there are from 1 to 10 [[NH₂]] amine groups for one episulfide group. After stirring at 60-100°C for from 0.2 to 3 hours, the solvent is distilled off under reduced pressure. To isolate the polymercaptopolyamine of formula [[I]] Ia or Ib, the excess amine reagent is removed by means of vacuum distillation at elevated temperature. In one embodiment of the invention, the excess amine is not removed and the mixture of the excess amine [[R₁-NH₂]] and the polymercaptopolyamine of formula [[I]] Ia or Ib is used as a hardener for epoxy resins.

Please replace the paragraph bridging pages 13 and 14 with the following paragraph:

In accordance with the above-mentioned procedure,

polymercaptopolyamines according to the invention (Examples I.1-I.19)

(Examples I.1-I.46) are prepared from the following diamines R₁ NH₂

amines and epoxides of formula H epoxy compounds of formula IIa or IIb:

BA: n-butylamine

OA: n-octylamine

CYA: cyclohexylamine

BZA: benzylamine

MBA: methylbutylamine

DMDP: N,H-demethyl-1,3-diaminopropane

MDP: N-methyl-1,3-diaminopropane

DACY: 1,2-diaminocyclohexane

AEP: N-2-aminoethylpiperazine

DETA: diethylenetriamine

IPD: isophoronediamine

MXDA: meta-xylylenediamine

DYTEK-A: 1,5-diamino-2-methylpentane

NBDA: isomeric mixture of 2,5- and 2,6-bis(aminomethyl)norbornane

NBDA: isomeric mixture of 2,3- and 2,0-bis(ammonicary), note the state of 2,5- and 2,0-bis(ammonicary), note that the state of 2,5- and 2,5- and

epoxide 1: ethylenediamine epoxide 1: liquid diglycidyl ether of bisphenol A having an epoxy

content of 5.25-5.4 eq./kg

epoxide 2: liquid mixture of diglycidyl ether of bisphenol A and

diglycidyl ether of bisphenol F having an epoxy content of

5.5-5.8 eq./kg

epoxide 3: 1,4-bis(hydroxymethyl)cyclohexane diglycidyl ether

epoxide 4: epoxy phenol novolak having an epoxy content of 5.6-5.8

eq./kg

epoxide 5: diglycidyl ether of hydrogenated bisphenol A

epoxide 6: tetraglycidyl ether of formula

epoxide 7: di(\(\mathcal{B}\)-methylglycidyl) ether of bisphenol A

epoxide 8: trimethylolpropane triglycidyl ether

epoxide 9: hexahydrophthalic acid diglycidyl ester (epoxy content:

5.6-6.2 eq./kg)

epoxide 10: phenyl glycidyl ether

Please replace the first footnote on page 16 with the following footnote:

*) mixture of amine [[R_1 -NH $_2$]] and polymercaptopolyamine; excess [[R_1 -NH $_2$]] amine not distilled off